METHOD 7190

CHROMIUM (ATOMIC ABSORPTION, DIRECT ASPIRATION)

- 1.0 SCOPE AND APPLICATION
 - 1.1 See Section 1.0 of Method 7000.
- 2.0 SUMMARY OF METHOD
 - 2.1 See Section 2.0 of Method 7000.
- 3.0 INTERFERENCES
 - 3.1 See Section 3.0 of Method 7000 if interferences are suspected.
- 3.2 An ionization interference may occur if the samples have a significantly higher alkali metal content than the standards. If this interference is encountered, an ionization suppressant (KCl) should be added to both samples and standards.
- 3.3 Background correction may be required because nonspecific absorption and scattering can be significant at the analytical wavelength. Background correction with certain instruments may be difficult at this wavelength due to low-intensity output from hydrogen or deuterium lamps. Consult the specific instrument manufacturer's literature for details.
- 4.0 APPARATUS AND MATERIALS
 - 4.1 For basic apparatus, see Section 4.0 of Method 7000.
 - 4.2 <u>Instrument parameters</u> (general):
 - 4.2.1 Chromium hollow cathode lamp.
 - 4.2.2 Wavelength: 357.9 nm.
 - 4.2.3 Fuel: Acetylene.
 - 4.2.4 Oxidant: Nitrous oxide.
 - 4.2.5 Type of flame: Fuel rich.
 - 4.2.6 Background correction: Not required.
- 5.0 REAGENTS
 - 5.1 See Section 5.0 of Method 7000.

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5.2 <u>Preparation of standards</u>:

- 5.2.1 Stock solution: Dissolve 1.923 g of chromium trioxide (CrO $_3$, analytical reagent grade) in Type II water, acidify with redistilled HNO $_3$, and dilute to 1 liter. Alternatively, procure a certified standard from a supplier and verify by comparison with a second standard.
- 5.2.2 Prepare dilutions of the stock solution to be used as calibration standards at the time of analysis. The calibration standards should be prepared using the same type of acid and at the same concentration as will result in the sample to be analyzed after processing.

6.0 SAMPLE COLLECTION. PRESERVATION. AND HANDLING

6.1 See Chapter Three, Section 3.1.3, Sample Handling and Preservation.

7.0 PROCEDURE

- 7.1 <u>Sample preparation</u>: The procedures for preparation of the sample are given in Chapter Three, Section 3.2.
 - 7.2 See Method 7000, Paragraph 7.2, Direct Aspiration.

8.0 QUALITY CONTROL

8.1 See Section 8.0 of Method 7000.

9.0 METHOD PERFORMANCE

9.1 The performance characteristics for an aqueous sample free of interferences are:

Optimum concentration range: 0.5-10 mg/L with a wavelength of 357.9 nm. Sensitivity: 0.25 mg/L. Detection limit: 0.05 mg/L.

- 9.2 For concentrations of chromium below 0.2 mg/L, the furnace procedure (Method 7191) is recommended.
- 9.3 Precision and accuracy data are available in Method 218.1 of Methods for Chemical Analysis of Water and Wastes.
- 9.4 The data shown in Table 1 were obtained from records of state and contractor laboratories. The data are intended to show the precision of the combined sample preparation and analysis method.

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10.0 REFERENCES

- 1. Methods for Chemical Analysis of Water and Wastes, EPA-600/4-82-055, December 1982, Method 218.1.
- 2. Gaskill, A., Compilation and Evaluation of RCRA Method Performance Data, Work Assignment No. 2, EPA Contract No. 68-01-7075, September 1986.

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TABLE 1. METHOD PERFORMANCE DATA

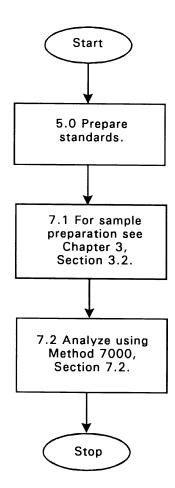
Sample Matrix	Preparation Method	Laboratory Replicates
Wastewater treatment sludge	3050	6,100, 6,000 ug/g
Emission control dust	3050	2.0, 2.8 ug/g

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